

(19)



European Patentamt
European Patent Office
Office européen des brevets



(11)

EP 0 971 404 A1

(12)

EUROPEAN PATENT APPLICATION

(43) Date of publication:
12.01.2000 Bulletin 2000/02

(51) Int. Cl.⁷: H01L 21/84, H01L 21/336,
H01L 27/12

(21) Application number: 99301978.5

(22) Date of filing: 15.03.1999

(84) Designated Contracting States:
AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU
MC NL PT SE
Designated Extension States:
AL LT LV MK RO SI

(30) Priority: 10.07.1998 US 113667

(71) Applicants:
• Sharp Kabushiki Kaisha
Osaka-shi, Osaka 545-8522 (JP)
• SHARP MICROELECTRONICS TECHNOLOGY,
INC.
Camas, WA 98607 (US)

(72) Inventors:
• Hsu, Sheng Teng
Camas, WA 98607 (US)
• Maa, Jer-Shen
Vancouver, WA 98684 (US)

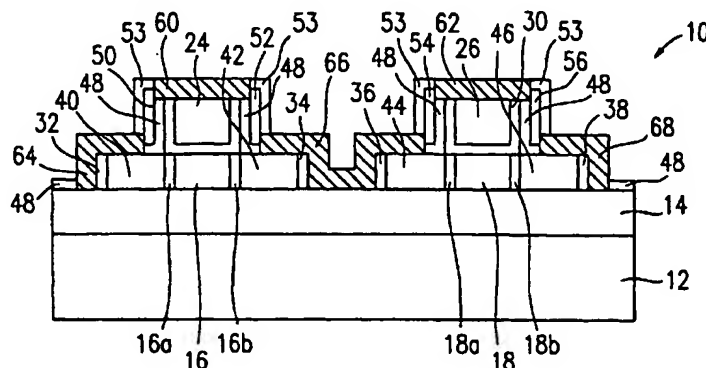
(74) Representative:
Brown, Kenneth Richard et al
R.G.C. Jenkins & Co.
26 Caxton Street
London SW1H 0RJ (GB)

(54) Double sidewall raised silicided source/drain CMOS transistor

(57) A method of forming a silicided device includes preparing a substrate by forming device areas thereon; providing structures that are located between the substrate and any silicide layers; forming a first layer of a first reactive material over the formed structures; providing insulating regions in selected portions of the structure; forming a second layer of a second reactive

material over the insulating regions and the first layer of first reactive material; reacting the first and second reactive materials to form silicide layers; removing any un-reacted reactive material; forming structures that are located on the silicide layers; and metallizing the device.

FIG. 5



EP 0 971 404 A1

Description

Field of the Invention

[0001] This invention relates to high performance CMOS formed on SIMOX and MOS transistors having very short channel length with shallow source and drain regions.

Background of the Invention

[0002] MOS circuits generally use a refractory metal, or silicide of a refractory metal, as a barrier, a conducting media, or an intermediate layer. Refractory metals and their suicides have relative low resistivities and low contact resistances and are desirable as conducting films and layers. Known silicide processes, however, fail to work on deep sub-micron MOS transistors because such processes generally consumes too much silicon. Additionally, impurities and problems achieving uniform deposition of silicide layers create manufacturing problems. Selective epitaxial deposition of silicon or selective deposition of polysilicon requires specialized manufacturing equipment. In addition, the selectivity of the silicide process is strongly dependant on the surface condition of the annealed film.

Summary of the Invention

[0003] The method of the invention for forming a silicided device includes preparing a substrate by forming device areas thereon; providing structures that are located between the substrate and any silicide layers; forming a first layer of a first reactive material over the formed structures; providing insulating regions in selected portions of the structure; forming a second layer of a second reactive material over the insulating regions and the first layer of first reactive material; reacting the first and second reactive materials to form silicide layers; removing any unreacted reactive material; forming structures that are located on the suicide layers; and metallizing the device.

[0004] It is an object of this invention to develop a simple, reliable, and cost effective silicide CMOS process/structure for very high density very small geometry circuit fabrication.

Brief Description of the Drawings

[0005]

Fig. 1 is a sectional front elevation of the structure following initial wafer preparation and LDD implantation.

Fig. 2 is a sectional front elevation of the structure following formation of N⁺ and P⁺ regions.

Fig. 3 is a sectional front elevation of the structure following deposition of a refractory metal layer.

Fig. 4 is a sectional front elevation of the structure following etching of the refractory metal layer.

Fig. 5 is a sectional front elevation of the structure following silicidation.

Fig. 6 is a sectional front elevation of the structure following selective etching of un-reacted refractory metal.

Fig. 7 is a sectional front elevation of the structure following selective etching of oxide and polysilicon layers.

Fig. 8 is a sectional front elevation of the completed structure.

Detailed Description of the Preferred Embodiments

[0006] The structure and the process of fabricating the structure according to the invention will be described using a SIMOX (Separation by IMplantation of Oxygen) substrate. The same technique may be applied to bulk silicon devices.

[0007] The starting material is a SIMOX wafer with very thin superficial silicon film. Referring now to Fig. 1, a portion of a SIMOX wafer is depicted generally at 10. Wafer 10 has a single crystal silicon portion 12, also referred to herein as the substrate. Buried oxide layer 14 has a thickness of between 100 nm and 300 nm, and the silicon film layer has a thickness not greater than 100 nm. The wafer is prepared to form device areas thereon. The structure is treated by active area etching, and threshold voltage adjustment ion implantation. In the case where bulk silicon is used, well diffusion is used, followed by LOCOS or proper isolation formation, threshold voltage adjustment, and ion implantation. In either case, the next step is gate oxidation, polysilicon deposition, gate electrode etching, and LDD ion implantation, to form those structures which are located between the substrate and any silicide layer.

[0008] The structure is sketched in Fig 1, and includes the substrate 12, a buried oxide layer 14, and two silicon regions 16, 18, which are the remnants of the superficial silicon layer. Portions of each silicon region 16, 18 are doped to form N⁺ regions 16a, 16b, and P⁺ regions 18a, 18b, respectively, with the central portion of each region remaining as untreated silicon. The doping density of regions 16 and 18 are $1.0 \cdot 10^{16} \text{ cm}^{-3}$ to $1.0 \cdot 10^{18} \text{ cm}^{-3}$ of boron and $5.0 \cdot 10^{15} \text{ cm}^{-3}$ to $5.0 \cdot 10^{17} \text{ cm}^{-3}$ of boron, respectively. The doping density of the N⁺ regions is $1.0 \cdot 10^{18} \text{ cm}^{-3}$ to $5.0 \cdot 10^{19} \text{ cm}^{-3}$ of As or phosphorous. The doping density of the P⁺ regions is $1.0 \cdot 10^{18} \text{ cm}^{-3}$ to $5.0 \cdot 10^{19} \text{ cm}^{-3}$ of boron. Silicon regions 16, 18 are surrounded by oxide caps 20, 22, respectively. Gate polysilicon regions 24, 26 are located above silicon regions 16, 18, respectively. The preceding steps may be achieved with any state-of-the-art process.

[0009] A layer of silicon oxide or silicon nitride is deposited, which layer functions as an insulator, over the entire substrate. The thickness of this insulating layer is between 50 nm to 100 nm. In the embodiment

described herein, silicon oxide is used. The structure is plasma etched, and now referring to Fig. 2, to remove the upper portions of the insulating layer oxide layer, leaving oxide at the sidewall of gate electrodes 24, 26, which, combined with the remains of oxide caps 20, 22, forms oxide cups 28, 30, and oxide sidewalls, 32, 34, 36 and 38 at the ends of silicon regions 16, 18.

[0010] A portion of the structure is covered with photoresist for N⁺ and P⁺ source/drain ion implantation for the nMOS and pMOS, respectively. N⁺ and P⁺ source/drain ions, i.e., As ions for the N⁺ regions and BF₂ ions for the P⁺ regions, are implanted at an energy level of 10 keV to 60 keV and a dose of $1.0 \cdot 10^{15} \text{ cm}^{-2}$ to $5 \cdot 10^{15} \text{ cm}^{-2}$ for the N⁺ region, and an energy level of 10 keV to 60 keV and a dose of $1 \cdot 10^{15} \text{ cm}^{-2}$ to $5.0 \cdot 10^{15} \text{ cm}^{-2}$ for the P⁺ region, forming N⁺ regions 40, 42 and P⁺ regions 44, 46, which will ultimately become the source/drain regions of the devices. The gate polysilicon prevents implantation of ions in the area directly beneath the gate polysilicon, which remains in their original state as silicon regions 16, 18. Silicon regions 16 and 18 are LDD regions, while regions 40, 46 are source regions and regions 42, 44 are drain regions.

[0011] Referring now to Fig. 3, a first layer 48 of a first reactive material is deposited over the already formed structures, followed by the formation of insulating regions 50, 52, 54 and 56 in selective portions of the structure, and the deposition of a second layer 58 of a second reactive material. In the first embodiment, first layer 48 is a thin layer of polysilicon, which is deposited over the entire structure to a thickness of between 50 nm to 100 nm. A layer of silicon oxide or silicon nitride is deposited to form insulating regions to a thickness of between 50 nm to 100 nm. Alternately, the oxide layer may be formed by a thermal process, to a thickness of 10 nm to 50 nm. The oxide or nitride layer is plasma etched to form oxide or nitride strips 50, 52, 54 and 56 at the sidewalls of gate electrode 24, 26, respectively. Second layer 58 is formed of a thin layer of refractory metal, which is deposited by CVD or sputtering. The refractory metal may be Co, Ti, Ni, and Pt, and is deposited to a thickness of between 5 nm and 50 nm.

[0012] The structure is covered with photoresist, and the refractory metal is etched out of the areas which will not have suicide located therein, as shown in Fig. 4. Silicidation takes place as a reaction between the refractory metal and the silicon during rapid thermal annealing (RTA) at a temperature of between 500 °C to 900 °C for 10 to 50 second, resulting in the formation of suicide layers 60, 62, 64, 66 and 68, as shown in Fig. 5.

[0013] The un-reacted refractory metal is removed by selective etching, with a solution such as $\text{NH}_4\text{OH} + \text{H}_2\text{O}_2 + \text{H}_2\text{O}$ for Ti, $\text{HNO}_3 + \text{HCl}$ for Pt and $\text{HCl} + \text{H}_2\text{O}_2$ for Ni or Co, resulting in the configuration shown in Fig. 6.

[0014] The remaining oxide is selective etched in a diluted BHF solution, and the polysilicon is selectively etched in an $\text{HNO}_3 : \text{H}_2\text{O}_2 : \text{H}_2\text{O}$ solution, resulting in the

configuration shown in Fig. 7. It should be noted that suicide layers 60, 62, located on the top of gate polysilicon 24, 26 has an overhang. Because the thickness of the polysilicon is no thicker than 100 nm, the overhang is less than 100 nm. Therefore, there is, with proper quality control in the manufacturing process, no step coverage problem.

[0015] Follow the state of the art process to complete the device fabrication to form any structures which are located on a suicide layer, above, or along side of a suicide layer, and which has not already been formed. The structure is covered with oxide 70 by CVD to a thickness of between 400 nm and 600 nm. Oxide layer 70 joins with oxide cups 28, 30. The structure is etched to form bores for metallization, and metal is deposited to form source electrode 72, gate electrode 74, combined drain electrode 76, gate electrode 78 and source electrode 80. A cross-sectional view of the finished CMOS pair is shown in Fig. 8.

[0016] In an alternate form of the invention, the refractory metal is deposited as the first reactive layer, the sidewall insulators formed, and a layer of polysilicon deposited as the second reactive layer. Portions of the second reactive layer, polysilicon in this case, is selectively etched, as in Fig. 4. Silicidation follows, and then selective etching of polysilicon and the refractory metal.

[0017] If the refractory metal is Ni, Co or Pt, a thin layer of Ti may be deposited on top of the initial metal layer. The thickness of Ti layer may be very thin such as 5 nm to 20 nm. The wafer is then exposed to air to convert Ti to titanium oxide. If necessary, the wafer is heated to a temperature 40 °C to 250 °C to convert all Ti to titanium oxide. The titanium oxide is plasma etched to form a titanium oxide sidewall at the side wall of the gate electrode. Polysilicon is deposited, photoresist is applied, and the polysilicon is etched out of the area where no silicide is needed. The wafer is then treated to form the silicide layers.

[0018] Although a preferred embodiment of the invention, and several variations thereof have been disclosed, it will be appreciated that further modifications and variations may be made thereto within the scope of the invention as defined in the appended claims.

Claims

1. A method of forming a silicided device, comprising:

- preparing a substrate by forming device areas thereon;
- providing structures that are located between the substrate and any suicide layers;
- forming a first layer of a first reactive material over the formed structures,
- providing insulating regions in selected portions of die structure;
- forming a second layer of a second reactive material over the insulating regions and the first

- layer of first reactive material;
 reacting the first and second reactive materials
 to form silicide layers;
 removing any un-reacted reactive material;
 forming structures that are located on the sili-
 cide layers; and
 metallizing the device.
2. The method of claim 1 wherein said reacting
 includes rapid thermally annealing the structure at
 a temperature of between about 500°C to 900 °C
 for a period of about 10 seconds to 50 seconds.
3. The method of claim 1 wherein said forming a first
 layer of a first reactive material over the formed
 structures includes depositing a layer of polysilicon
 and wherein said forming a second layer of a sec-
 ond reactive material includes depositing a layer of
 a refractory metal taken from the group of refractory
 metals consisting of Ni, Co, Ti and Pt.
4. The method of claim 1 wherein said forming a first
 layer of a first reactive material over the formed
 structures includes depositing a layer of a refractory
 metal taken from the group of refractory metals
 consisting of Ni, Co, Ti and Pt, and wherein said
 forming a second layer of a second reactive mate-
 rial includes depositing a layer of polysilicon.
5. The method of claim 1 wherein said forming a first
 layer of a first reactive material over the formed
 structures includes depositing a layer of a refractory
 metal taken from the group of refractory metals
 consisting of Ni, Co and Pt, and which includes
 depositing a layer of Ti on top of the first layer of first
 reactive material; and wherein said providing insu-
 lating regions in selected portions of the structure
 includes oxidizing the Ti layer to form TiO₂; and
 wherein said forming a second layer of a second
 reactive material includes depositing a layer of poly-
 silicon.
6. A method of forming a silicided device, comprising:
 preparing a substrate by forming device areas
 thereon;
 providing structures that are located between
 the substrate and any silicide layers;
 depositing a layer of polysilicon over the
 formed structures;
 providing insulating regions in selected por-
 tions of the structure; depositing a layer of a
 refractory metal taken from the group of refrac-
 tory metals consisting of Ni, Co, Ti and Pt over
 the insulating regions and the polysilicon layer;
 reacting the polysilicon and the refractory
 metal to form silicide layers;
 removing any un-reacted refractory metal;
- forming structures that are located on the sili-
 cide layers; and
 metallizing the device.
7. The method of claim 6 wherein said reacting
 includes rapid thermally annealing the structure at
 a temperature of between about 500°C to 900°C for
 a period of about 10 seconds to 50 seconds.
8. A method of forming a silicided device, comprising:
 preparing a substrate by forming device areas
 thereon;
 providing structures that are located between
 the substrate and any silicide layers;
 depositing a layer of a refractory metal over the
 formed structures;
 providing insulating regions in selected por-
 tions of the structure;
 depositing a layer of polysilicon over the insu-
 lating regions and the first layer of first reactive
 material;
 reacting the first and second reactive materials
 to form silicide layers;
 removing any un-reacted reactive material;
 forming structures that are located on the sili-
 cide layers; and
 metallizing the device.
9. The method of claim 8 wherein said reacting
 includes rapid thermally annealing the structure at
 a temperature of between about 500°C to 900°C for
 a period of about 10 seconds to 50 seconds.
10. The method of claim 8 wherein said depositing a
 layer of a refractory metal taken from the group of
 refractory metals consisting of Co and Pt, and
 which includes depositing a layer of Ti on top of the
 first layer of first reactive material; and wherein said
 providing insulating regions in selected portions of
 the structure includes oxidizing the Ti layer to form
 TiO₂; and wherein said forming a second layer of a
 second reactive material includes depositing a layer
 of polysilicon.
11. The method of claim 8 wherein said depositing a
 layer of a refractory metal includes depositing a
 refractory metal taken from the group of refractory
 metals consisting of Ni, Co, Ti and Pt.
12. A method of forming silicide portions (60, 62, 64,
 66, 68) on a substrate (12, 14) bearing formed
 devices, wherein two layers of reactive materials
 (48, 58) are deposited, together with insulating por-
 tions (50, 52, 54, 56) sandwiched between said lay-
 ers (48, 58) in selected areas other than those of
 the required suicide portions, and wherein the lay-
 ers are reacted and the unreacted portions thereof

are removed along with the insulating portions
sandwiched therebetween.

5

10

15

20

25

30

35

40

45

50

55

5

FIG. 1

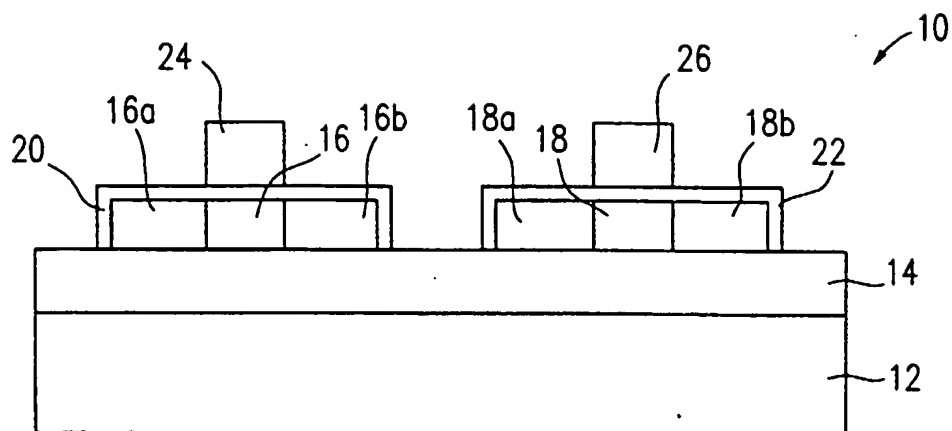


FIG. 2

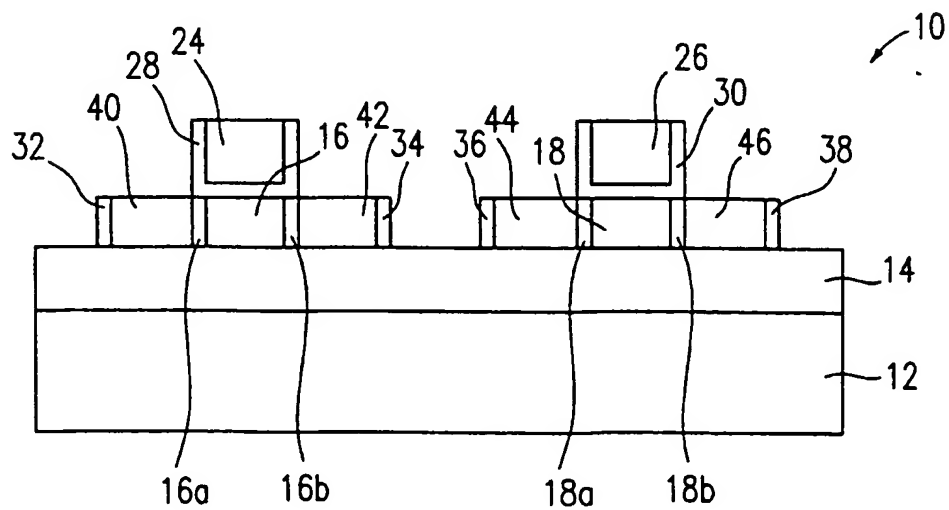


FIG. 3

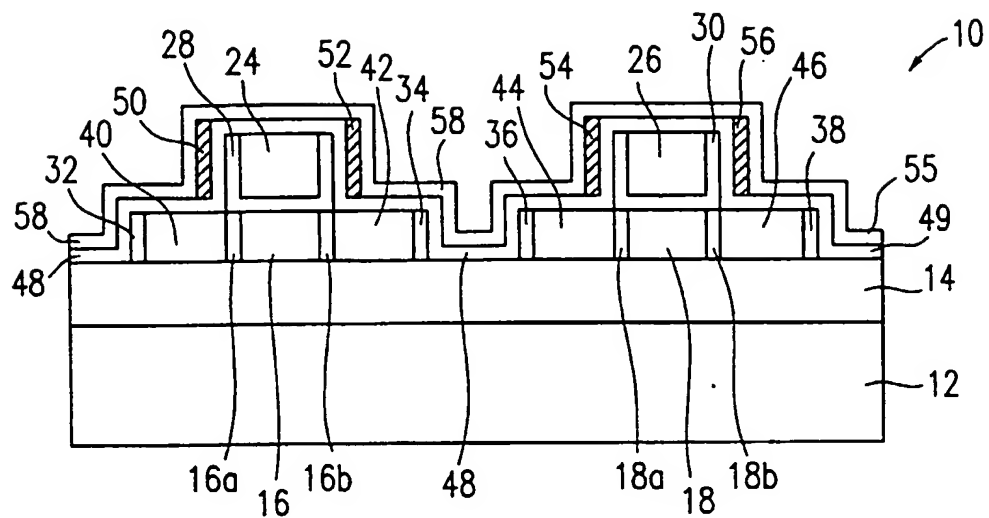


FIG. 4

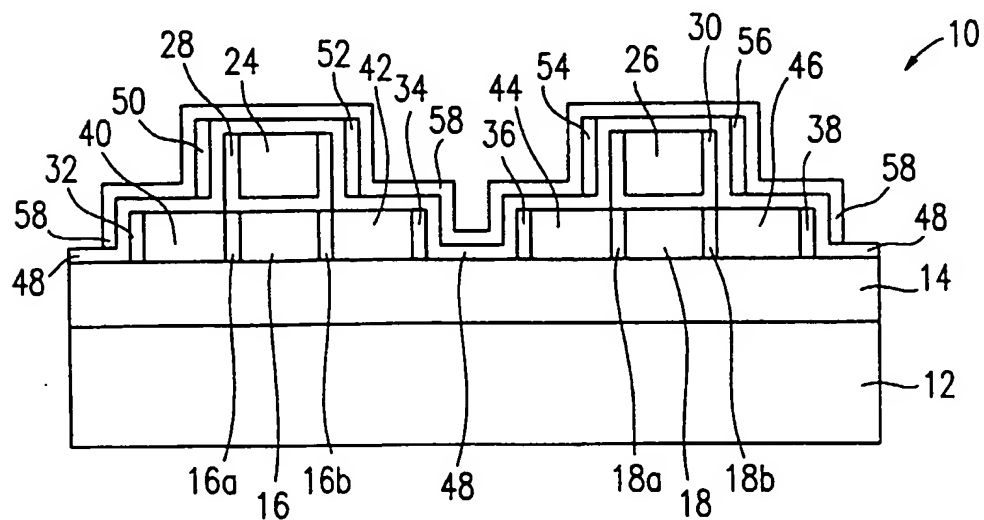


FIG. 5

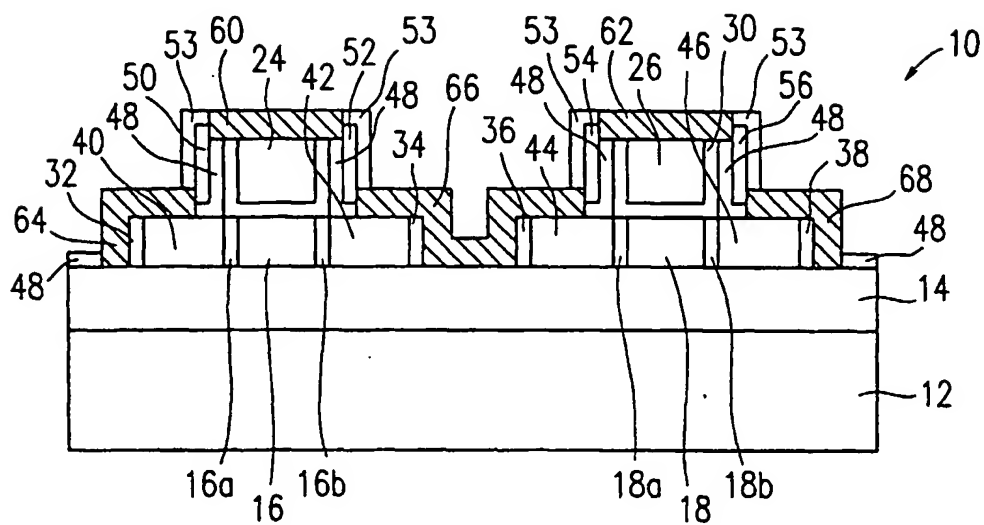


FIG. 6

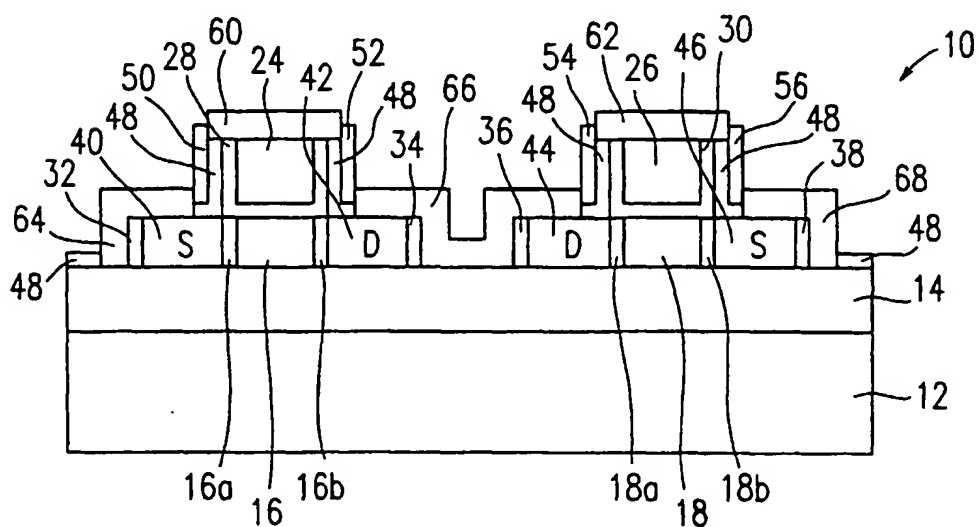


FIG. 7

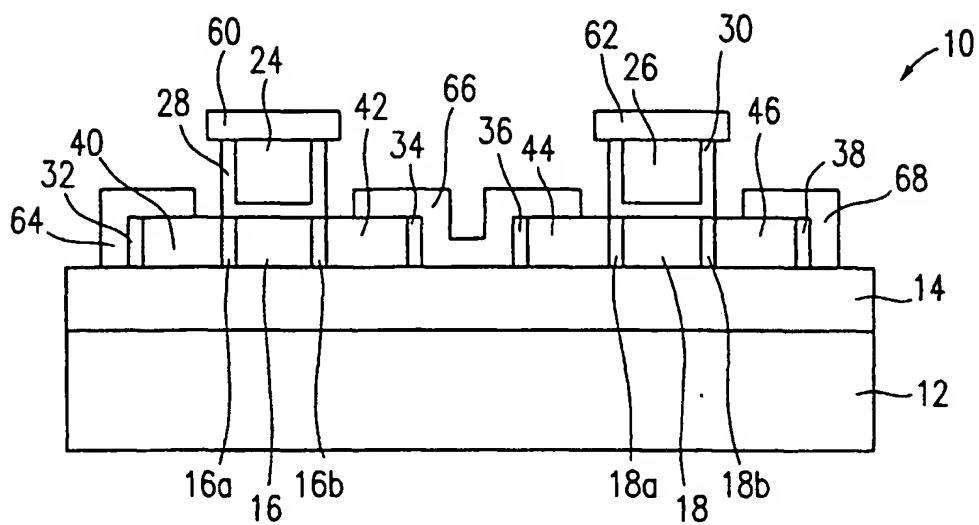
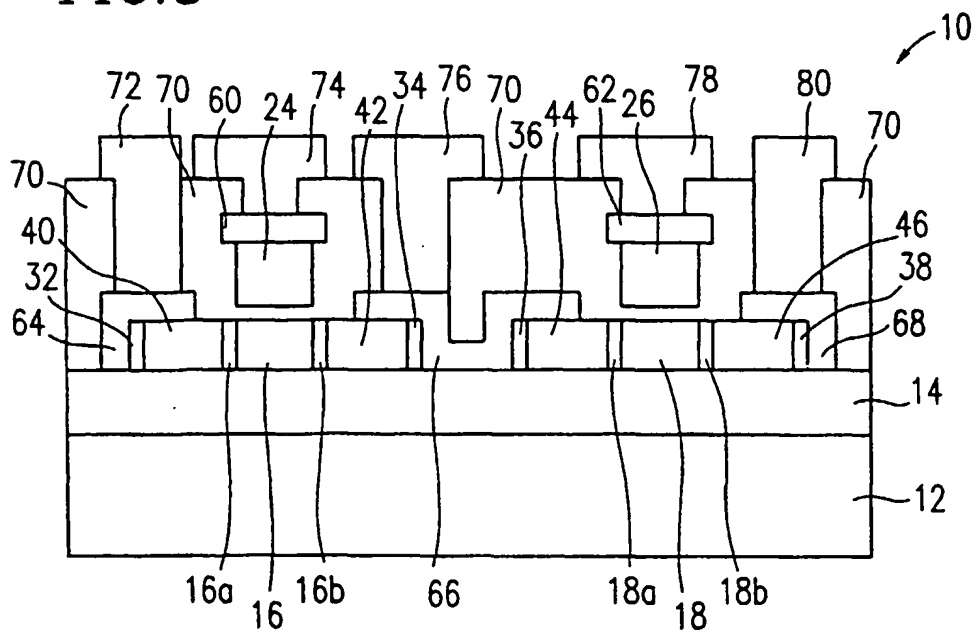


FIG. 8





European Patent
Office

EUROPEAN SEARCH REPORT

Application Number
EP 99 30 1978

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.7)
X	EP 0 766 305 A (SIEMENS AG) 2 April 1997 (1997-04-02)	1,4,6,8, 11,12	H01L21/84 H01L21/336
Y	* the whole document *	2,7,9	H01L27/12
A	---	5,10	
Y	EP 0 480 446 A (TEXAS INSTRUMENTS INC) 15 April 1992 (1992-04-15)	2,7,9	
A	* page 5, column 7, line 6 - page 14, column 25, line 9; figures 1-19 *	1,3,4,6, 8,11,12	
X	US 5 464 782 A (KOH CHAO-MING) 7 November 1995 (1995-11-07)	1-4,6-9, 11	
A	* the whole document *	5,10,12	
A	US 5 365 111 A (RAMASWAMI SESHADRI ET AL) 15 November 1994 (1994-11-15)	1-4,6-9, 11,12	
	* column 2, line 40 - column 4, line 68; figures 1-3; table 1 *		
A	PATENT ABSTRACTS OF JAPAN vol. 1995, no. 04, 31 May 1995 (1995-05-31)	1,3,4,6, 8,12	TECHNICAL FIELDS SEARCHED (Int.Cl.7)
	-& JP 07 030104 A (TOSHIBA CORP), 31 January 1995 (1995-01-31)		H01L
	* abstract *		
P,X	WO 98 35380 A (ADVANCED MICRO DEVICES INC) 13 August 1998 (1998-08-13)	1-4,6-9, 11	
P,A	* the whole document *	12	

The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 15 October 1999	Examiner Albrecht, C
CATEGORY OF CITED DOCUMENTS		T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document	
X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document			

EPO FORM 1503 03.82 (Pct/Co1)

**ANNEX TO THE EUROPEAN SEARCH REPORT
ON EUROPEAN PATENT APPLICATION NO.**

EP 99 30 1978

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report. The members are as contained in the European Patent Office EDP file on
The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

15-10-1999

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
EP 0766305 A	02-04-1997	DE 19536249 A	10-04-1997
		JP 9134971 A	20-05-1997
		US 5733803 A	31-03-1998
EP 0480446 A	15-04-1992	US 5168072 A	01-12-1992
		JP 6077246 A	18-03-1994
		US 5397909 A	14-03-1995
US 5464782 A	07-11-1995	NONE	
US 5365111 A	15-11-1994	JP 6232073 A	19-08-1994
		US 5451545 A	19-09-1995
JP 07030104 A	31-01-1995	NONE	
WO 9835380 A	13-08-1998	NONE	

EPO FORM P0459

For more details about this annex : see Official Journal of the European Patent Office, No. 12/82